MEDISTIK ICE- menthol, camphor solution Natureteq Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

NATURETEQ - MEDISTIK PRO ICE THERAPY, 50231-411

ACTIVE INGREDIENTS

Menthol 10%

Camphor 4%

PURPOSE

Topical Analgesic

USES

For the temporary relief of sore muscles and joints associated with

- strains and sprains
- backaches
- lumbago
- pain of tendons and/or ligaments
- arthritic or rheumatic pain

WARNINGS

For external use only. Avoid contact with eyes and mucous membranes. Do not apply to wounds or damaged skin. Do not tightly bandage. Do not apply with external heat, such as an electric heating pad, as this may result in excessive skin irritation or skin burn.

Stop use and consult a physician if condition worsens, rash or irritation develops, or if symptoms persist for more than 7 days or clear up and recur in a few days. Consult a health care practitioner prior to use if you are pregnant or breastfeeding. Rashes and/or burning discomfort, and hypersensitivity such as allergy have been known to occur; in which case, discontinue use.

DIRECTIONS

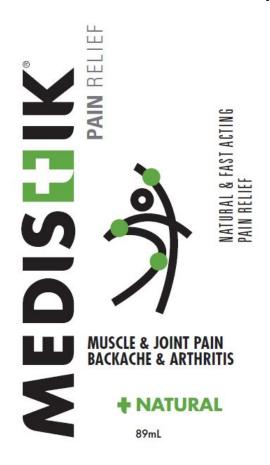
For use on adults and children over 12 years of age.

Apply to affected area(s) 3 to 4 times per day as required. A stinging or burning sensation will be experienced during the first few minutes as the formula begins working. For arthritis or muscle pain of the hands, retain for at least 10 minutes then wash hands.

INACTIVE INGREDIENTS

Denatured alcohol, Water, Peppermint oil, Glycerin, Acrylates copolymer, MSM, Ilex paraguariensis leaf extract, Glucosamine sulphate, Chondroitin sulphate, Folic acid, Vitamin C, Vitamin D.

Cautions: Keep out of reach of children. If overdose or accidental ingestion occurs, call a Poison Control Center immediately.





Drug Facts	
Active Ingredients Menthol 10% Camphor 4%	
Uses Temporarily relieves minor aches muscles and joints associated with ■ sin ■ arthritis ■ strains ■ sprains ■ bruises ■	nple backache
Warnings For external use only	

When using this product was only as directed avoid getting into the eyes and mucous membranes do not bandage tightly or use with a heating pad do not apply to wounds or damaged skin

Stop use and ask doctor if ■ condition worsens
■ symptoms persist for more than 7 days or clear up and occur again within a few days ■ excessive skin irritation or redness occurs

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children to avoid accidental ingestion. If swallowed, get medical help or contact a Poison Control Center immediately.

Flammable: Keep away from excessive heat or open flame. Store in a cool dry place with lid or cap closed tightly

Directions and adults and children 12 years of age and older: apply product on affected area, not more than 4 times daily a children under 12 years of age: ask a doctor

Inactive Ingredients ■ Denatured alcohol, Water, Peppermint oil, Glycerin, Polyacrylate Crosspolymer-6, MSM, Ilex paraguariensis leaf extract, Glucosamine sulfate, Chondroitin sulfate, Folic acid. Vitamin C. Vitamin D. may contain sodium hydroxide

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MEDISTIK ICE

menthol, camphor solution

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50231-411
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	10 g in 100 mL
CAMPHOR OIL (UNII: 75IZZ8Y727) (CAMPHOR OIL - UNII:75IZZ8Y727)	CAMPHOR OIL	4 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)	
GLUCOSAMINE SULFATE (UNII: 1FW7WLR731)	
CHONDROITIN SULFATE (CHICKEN) (UNII: 7VZ 9466BAB)	
FOLIC ACID (UNII: 935E97BOY8)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
VITAMIN D (UNII: 9VU1KI44GP)	

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50231-411- 11	89 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/21/2020	
2	NDC:50231-411- 12	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/18/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	02/17/2016	

Labeler - Natureteq Inc. (243737371)

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